(FILE 'HOME' ENTERED AT 15:33:14 ON 19 MAR 2003)

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SEA CD20(10W)B CELL

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FILE 'DGENE, BIOSIS, MEDLINE, SCISEARCH, EMBASE, CAPLUS, CANCERLIT, TOXCENTER, USPATFULL, ESBIOBASE, BIOTECHNO, PROMT, DRUGU, LIFESCI, ADISCTI, JICST-EPLUS, CIN, GENBANK, IFIPAT, PASCAL, PHARMAML, WPIDS, PHIN, CONFSCI, FEDRIP, DRUGNL, ADISNEWS, BIOCOMMERCE, ...' ENTERED AT 15:35:41 ON 19 MAR 2003

11897 S CD20(10W)B CELL

39 S VACCIN? (10W) CD20

L2 L3

L1

L4	18	DUP REM L3 (21 DUPLICATES REMOVED)
L5	42	S CD20(25W)(EXTRACELL?) AND DOMAIN
L6	30	DUP REM L5 (12 DUPLICATES REMOVED)
L7		S CD20(10W)ANTIBOD?
L8	17	S L7 AND CD10(10W) (ANTIGEN OR IMMUNOGEN?)
L9	16	DUP REM L8 (1 DUPLICATE REMOVED)
L10		S B1(20W)CD20 AND (B CELL)
L11	148	DUP REM L10 (224 DUPLICATES REMOVED)
L12		S L10 AND B1(25W)ANTIBOD?
L13	75	DUP REM L12 (91 DUPLICATES REMOVED)
L14	425	S B1(25W)VACCINE
L15	250	DUP REM L14 (175 DUPLICATES REMOVED)
L16		S L14 AND (CANCER OR TUMOR OR TUMOUR)
L17	15	DUP REM L16 (4 DUPLICATES REMOVED)

L4 ANSWER 16 OF 18 CAPLUS COPYRIGHT 2003 ACS

AN 1997:8270 CAPLUS

DN 126:103042

TI Lysis of syngeneic tumor B cells by autoreactive cytotoxic T lymphocytes specific for a CD19 antigen-derived synthetic peptide

AU Hooijberg, Erik; Visseren, Marjan J. W.; Van Den Berk, Paul C. M.; Jellema, Anke P.; Romeijn, Petra; Sein, Johan J.; Van Der Voort, Ellen I. H.; Hekman, Annemarie; Ossendorp, Ferry; Melief, Cornelis J. M.

CS Department of Immunology, The Netherlands Cancer Institute/Antoni van Leeuwenhoek Huis, Amsterdam, Neth.

Journal of Immunotherapy with Emphasis on Tumor Immunology (1996), 19(5), 346-356

CODEN: JIEIEZ; ISSN: 1067-5582

PB Lippincott-Raven

DT Journal

LA English

Cytotoxic T lymphocytes (CTL) play an important role in the destruction of AΒ immunogenic tumors. A novel category of target antigens for CTL concerns normal differentiation antigens as most clearly demonstrated in human melanoma. In the case of B-cell cancers, differentiation antigens normally expressed on B cells may be useful targets. In this report, we have focused on the murine B-cell differentiation antigens CD19 and CD20. We have identified 18 peptide sequences on the basis of major histocompatibility complex (MHC) class-I binding-motifs as candidates for the induction of autoreactive CTL. Six of the peptides were capable of binding efficiently to either Kb or Db and were subsequently used for in vivo induction of CTL. Vaccination with each of three peptides led to peptide-specific CTL. Two peptides were derived from the mCD20 antigen and one from the mCD19 antigen. CTL specific for the mCD19-derived peptide were also capable of killing a syngeneic B-cell tumor line. Recognition of the peptide as well as the tumor cells was shown to be Kb restricted. This is the first report to show that autoreactive CTL recognizing peptides derived from B-cell-specific differentiation antigens can be generated by vaccination with a synthetic peptide.

IT Lymphoma

(B-cell; vaccination with CD19 or CD20 peptides induces cytotoxic T-lymphocytes that lyse syngeneic B-cell tumors in mice)

IT 185856-50-2 185856-51-3

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(vaccination with CD19 or CD20 peptides induces

cytotoxic T-lymphocytes that lyse syngeneic B-cell tumors in mice)

Japan or John and